(26) Cosmetic product ingredient and cosmetic raw material composition statements, § 720.8 of this chapter.

(27) Cosmetic product experience reports, in § 730.7 of this chapter.

(28) Device premarket notification submissions, in § 807.95 of this chapter.

(29) Electronic product information, in §§ 1002.4 and 1002.42 of this chapter.

(30) Data and information submitted to the Commissioner or to classification panels in connection with the classification or reclassification of devices intended for human use, in § 860.5 of this chapter.

(31) Data and information submitted in offers to develop a proposed performance standard for medical devices, in § 861.26 of this chapter.

(32) Investigational device exemptions in § 812.38 of this chapter.

(33) Health claims petitions, in § 101.70 of this chapter.

(34) Premarket approval application, in § 814.9 of this chapter.

(35) Report of certain adverse experiences with a medical device, in § 803.9 of this chapter.

(36) Disqualification determination of an institutional review board, in § 56.122 of this chapter.

(37) Disqualification determination of a nonclinical laboratory, in § 58.213 of this chapter.

(38) Minutes or records regarding a public advisory committee, in § 14.65(c) of this chapter.

(39) Data submitted regarding persons receiving an implanted pacemaker device or lead, in § 805.25 of this chapter.

(40) Humanitarian device exemption application, in § 814.122 of this chapter.

(41) Premarket notifications for food contact substances, in § 170.102 of this chapter.

(42) Registration of food facilities, in § 1.243 of this chapter.

(43) Minor-use or minor-species (MUMS) drug designations, in § 516.52 of this chapter.

(44) Minor-species drug index listings, in § 516.171 of this chapter.

[42 FR 15616, Mar. 22, 1977, as amended at 42 FR 19989, Apr. 15, 1977; 42 FR 42526, Aug. 28, 1977; 42 FR 58889, Nov. 11, 1977; 43 FR 32993, July 28, 1978; 51 FR 22475, June 19, 1986; 54 FR 9038, Mar. 3, 1989; 58 FR 2533, Jan. 6, 1993; 59 FR 536, Jan. 5, 1994; 61 FR 33244, June 26, 1996; 62 FR 40592, July 29, 1997; 64 FR 56448, Oct. 20, 1999; 67 FR 13717, Mar. 26, 2002; 67 FR 35729, May 21, 2002; 68 FR 58965, Oct. 10, 2003; 72 FR 41017, July 26, 2007; 72 FR 69118, Dec. 6, 2007]

§ 20.101 Administrative enforcement records.

(a) All Food and Drug Administration records relating to administrative enforcement action disclosed to any member of the public, including the person who is the subject of such action, are available for public disclosure at the time such disclosure is first made. Such records include correspondence with companies following factory inspection, recall or detention requests, notice of refusal of admission of an imported product, regulatory letters, information letters, Forms FD–483 and FD–2275 furnished to companies after factory inspection, and similar records.

(b) To the extent that any of such records fall within the exemption for investigatory records established in § 20.64, the Commissioner determines that they are subject to discretionary release pursuant to § 20.82.

(c) Records relating to administrative enforcement action that are not disclosed to any member of the public constitute investigatory records that are subject to the rules for disclosure established in § 20.64. For example, an establishment inspection report is an investigatory record and thus subject to § 20.64 except insofar as the Commissioner exercises his discretion to release it pursuant to § 20.82.

§ 20.102 Court enforcement records.

(a) All records and documents filed in the courts are available for public disclosure unless the court orders otherwise. The Food and Drug Administration will make available for public disclosure such records or documents if the agency can determine that it has an accurate copy of the actual record or document filed in the court. If the Food and Drug Administration cannot